## **CLAIMS**

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- 1. Use of a biological material containing:
- c) a three-dimensional matrix based on a hyaluronic acid derivative and optionally
- d) chondrocytes and/or mesenchymal cells partially or completely differentiated towards chondrocytes,
- for the preparation of a graft to be surgically implanted into a joint cartilage damaged by or to be protected against a degenerative and/or inflammatory pathology, selected from osteoarthritis and/or osteoarthrosis, rheumatoid arthritis and psoriatic arthritis.
- 2. The use according to claim 1, wherein, when the biological material contains the aforementioned cellular components (b) said graft is an *in vitro* cartilage tissue to be surgically implanted *in vivo* inside the inflamed joint capsule in which one of said degenerative pathologies has been established with consequent degradation of the extracellular cartilage matrix.
- 3. The use according to claim 2 wherein *in vitro* cartilage tissue further comprise the extracellular matrix produced by said chondrocytes or mesenchymal cells partially or completely differentiated towards chondrocytes said extracellular matrix being both inside said *in vitro* cartilage tissue and once *in vivo* implanted also inside the joint cartilage affected by one of said degenerative pathologies.
- 4. The use according to anyone of claims 1-3, wherein said grafts are to be surgically into joint cartilage in the early stages of one of said degenerative diseases.
  - 5. The use according to claim 4 wherein said grafts are to be surgically implanted at the beginning of the process of degradation of the molecules that make up the extracellular matrix of the cartilage.
  - 6. The use according to anyone of claims 1-3 wherein said grafts are to be surgically implanted in the later stages of said pathology too, when moderately and/or badly damaged areas of cartilage can be seen.
- 7. The use according to anyone of claims wherein the average molecular weight of hyaluronic acid in the hyaluronic acid derivative range between 1x 10<sup>5</sup>Da and 1x 10<sup>6</sup>Da.
  - 8. The use according to claim 7, wherein the average molecular weight of

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hyaluronic acid range between 200,000 and 750,000 Da.

- 9. The use according to anyone of claims 1-8, wherein the hyaluronic acid derivative is selected from the class consisting of:
- A) HA salified with organic and/or inorganic bases,
- B) HA esters with alcohols of the aliphatic, araliphatic, cycloaliphatic, aromatic, cyclic and heterocyclic series,
  - C) HA esters with alcohols of the aliphatic, araliphatic, cycloaliphatic, aromatic, cyclic and heterocyclic series.
  - D) O-sulphated derivatives of HA,
- E) inner esters of HA with a percentage of esterification that does not exceed 20%,
  - F) Deacetylated derivatives of HA obtained by the deacetylation of the N-acetylglucosamine fraction,
  - G) percarboxylated derivatives of HA obtained by oxidising the primary hydroxyl of the N-acetyl-glucosamine fraction with a degree of percarboxylation ranging between 0.1 and 100%.
  - 10. The use according to claim 9, wherein the HA derivative belongs to class(A) it is obtained by treating hyaluronic acid with sodium hydroxide.
- 11. The use according to claim 9, wherein, when the HA derivative belongs to class (B) it has a percentage of esterification ranging from 50 to 100%, and the remaining percentage of unesterified HA is salified with organic or inorganic base.
  - 12. The use according to claim 11, wherein said base is sodium hydroxide.
  - 13. The use according to claim 9, wherein when the HA derivative belongs to class
- (C) it has a percentage of amidation ranging between 0.1 and 50% and the remaining portion is salified with organic and/or inorganic bases.
  - 14. The use according to claim 13 wherein said base is sodium hydroxide.
  - 15. The use according to claim 9, wherein when the HA derivative belongs to class
  - (D) it has from 1 to 4 -OSO<sub>3</sub>H group per saccharide unit.
- 16. The use according to claim 9 wherein when the HA derivative belongs to class (E) it has a degree of esterification ranging from 0.05 to 10%, and the remaining percentage of non-esterified HA may be salified with organic and/or inorganic

## bases.

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- 17. The use according to claim 16, wherein said base is sodium hydroxide.
- 18. The use according to claim 9, wherein when the HA derivative belongs to class
- (E) it has a percentage of deacetylation ranging between 0.1 and 30% and all the carboxy groups of HA are salified with organic and/or inorganic bases.
- 19. The use according to claim 18, wherein said base is sodium hydroxide.
- 20. The use according to claim 9, wherein, when the HA derivative belongs to class
- (G), it has a degree of percarboxylation ranging from 25 to 75% and all the carboxy groups are salified with organic and/or inorganic bases.
- 10 21. The use according to claim 20, wherein the base is sodium hydroxide.
  - 22. The use according to anyone of claims 1-21, wherein said three-dimensional matrix is in a form selected from the group consisting of: a non-woven tissue, a tissue, microspheres, and a sponge.
  - 23. The use according to anyone of claims 1-22, wherein said HA derivative is a hyaluronic acid ester belonging to class (A).
  - 24. The use according to claim 23 wherein said HA ester is the benzyl ester having a percentage of esterification ranging from 75 to 100%.
  - 25. The use according to claim 24, wherein said benzylester has a percentage of esterification of 100% and is in the form of a non-woven tissue.